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Time: 14:09:22

Inventor Information for 10/668475

Inventor Name	City	State/Country					
DOLLE, ROLAND E.	KING OF PRUSSIA	PENNSYLVANIA					
HOYER, DENTON W.	EXTON	PENNSYLVANIA					
ROSS, TINA MORGAN	AUDUBON	PENNSYLVANIA					
RINKER, JAMES M.	READING	PENNSYLVANIA					
SCHMIDT, STANLEY J.	CHESTER SPRINGS	PENNSYLVANIA					
ATOR, MARK A.	PAOLI	PENNSYLVANIA					
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NEWS 20 JAN 27 Source of Registration (SR) information in REGISTRY updated
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NEWS 21 JAN 27 A new search aid, the Company Name Thesaurus, available in
                 CA/CAplus
                 German (DE) application and patent publication number format
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             DECEMBER 28 CURRENT WINDOWS VERSION IS V7.00, CURRENT
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              MACINTOSH VERSION IS V6.0b(ENG) AND V6.0Jb(JP),
              AND CURRENT DISCOVER FILE IS DATED 23 SEPTEMBER 2003
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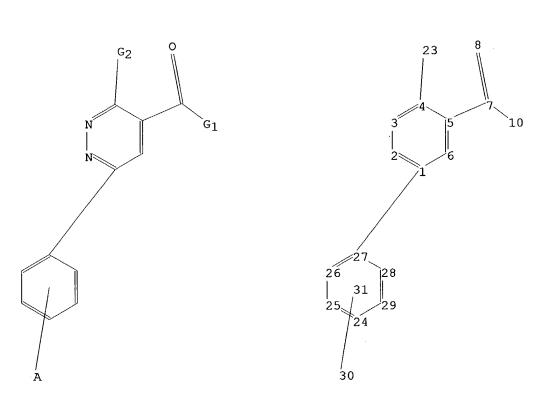
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chain nodes : 7 8 10 17 18 19 23 30 ring nodes : chain bonds : 1-27 4-23 5-7 7-8 7-10 14-17 18-19 ring bonds : $1-2^{-1}$ 1-6 2-3 3-4 4-5 5-6 11-12 11-16 12-13 13-14 14-15 15-16 24-2524-29 25-26 26-27 27-28 28-29 exact/norm bonds : 4-23 7-8 7-10 14-17 18-19 exact bonds : 1-27 5-7 normalized bonds : $1-2 \quad 1-6 \quad 2-3 \quad 3-4 \quad 4-5 \quad 5-6 \quad 11-12 \quad 11-16 \quad 12-13 \quad 13-14 \quad 14-15 \quad 15-16 \quad 24-25$ 24-29 25-26 26-27 27-28 28-29

G1:0,N

G2:X,S,P,[*1],[*2]

Match level:

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:CLASS 8:CLASS 10:CLASS 11:Atom 12:Atom 13:Atom 14:Atom 15:Atom 16:Atom 17:CLASS 18:CLASS 19:CLASS 23:CLASS 24:CLASS 25:Atom 26:Atom 27:Atom 28:Atom 29:Atom 30:CLASS 31:CLASS

L1 STRUCTURE UPLOADED

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SAMPLE SEARCH INITIATED 13:02:29 FILE 'REGISTRY' SAMPLE SCREEN SEARCH COMPLETED - 22 TO ITERATE

100.0% PROCESSED 22 ITERATIONS

2 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 159 TO 721 PROJECTED ANSWERS: 2 TO 124

L2 2 SEA SSS SAM L1

=> d scan

L2 2 ANSWERS REGISTRY COPYRIGHT 2004 ACS on STN

IN 4-Pyridazinecarboxamide, 3-chloro-6-methyl-5-(4-nitrophenyl)-N-phenyl-,
1-oxide (9CI)

MF C18 H13 C1 N4 O4

$$\begin{array}{c} \mathsf{R} \\ & \mathsf{C-}\,\mathsf{NHPh} \\ || \\ & \mathsf{O} \end{array}$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):.

L2 2 ANSWERS REGISTRY COPYRIGHT 2004 ACS on STN

IN 4-Pyridazinecarboxamide, 3-chloro-6-(4-cyanophenyl)- (9CI)

MF C12 H7 C1 N4 O

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

ALL ANSWERS HAVE BEEN SCANNED

=> s 11 full FULL SEARCH INITIATED 13:03:07 FILE 'REGISTRY' FULL SCREEN SEARCH COMPLETED - 421 TO ITERATE

100.0% PROCESSED 421 ITERATIONS SEARCH TIME: 00.00.01

117 ANSWERS

L3 117 SEA SSS FUL L1

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=> s 13

L4 2 L3

=> d 1-2 all

```
L4
     ANSWER 1 OF 2 CAOLD COPYRIGHT 2004 ACS on STN
     CA62:6480a CAOLD
AN
TΙ
     synthesis of N-containing heterocycles - (XXXII) syntheses of
     pyrazolo[3,4-c]-pyridazines
     Dornow, Alfred; Abele, W.
ΑU
                1430-22-4
                             1430-23-5
IT
     271-75-0
                                         1445-53-0
     1445-54-1
                1445-55-2
                            1445-56-3
                                         1933-44-4
                                                     2125-84-0
     2125-85-1
                 2125-86-2
                               2125-87-3
                                             2125-88-4
                                                     2125-93-1
                                                                 2125-94-2
                 2125-90-8
                            2125-91-9
                                         2125-92-0
     2125-89-5
                3052-82-2 88526-64-1
                                        88891-82-1 91113-85-8
     2283-61-6
    ANSWER 2 OF 2 CAOLD COPYRIGHT 2004 ACS on STN
L4
AN
     CA53:22026b CAOLD
     6-chloropyridazines
TΙ
     CIBA Ltd.
PA
DT
     Patent
     PATENT NO.
                  KIND
                                DATE
PΙ
     GB 807548
     7083-49-0 54108-27-9 93824-72-7 94011-79-7
IT
     101884-08-6 107919-08-4
=> d 1-2 all hitstr
    ANSWER 1 OF 2 CAOLD COPYRIGHT 2004 ACS on STN
T.4
    CA62:6480a CAOLD
AN
TΙ
     synthesis of N-containing heterocycles - (XXXII) syntheses of
     pyrazolo[3,4-c]-pyridazines
     Dornow, Alfred; Abele, W.
ΑU
                           1430-23-5
     271-75-0 1430-22-4
IT
                                         1445-53-0
                                                     2125-84-0
                            1445-56-3
                                         1933-44-4
     1445-54-1
               1445-55-2
     2125-85-1
               2125-86-2
                               2125-87-3
                                             2125-88-4
                                                     2125-93-1
     2125-89-5
                2125-90-8
                            2125-91-9
                                         2125-92-0
                                                                 2125-94-2
                 3052-82-2 88526-64-1 88891-82-1 91113-85-8
     2283-61-6
IT
     1445-53-0
                   1445-54-1
                                 2125-86-2
     2125-87-3
                   2125-88-4
RN
     1445-53-0 CAOLD
     4-Pyridazinecarboxylic acid, 3-chloro-6-methyl-, ethyl ester (7CI, 8CI)
CN
     (CA INDEX NAME)
```

CN 4-Pyridazinecarboxylic acid, 3-chloro-, ethyl ester (7CI, 8CI, 9CI) (CA INDEX NAME)

RN 2125-86-2 CAOLD

CN 4-Pyridazinecarboxylic acid, 3-chloro-6-methyl-, hydrazide (7CI, 8CI) (CA INDEX NAME)

$$\begin{array}{c|c} & & \text{C1} \\ & & \\ \text{Me} & & \\ & &$$

RN 2125-87-3 CAOLD

CN 4-Pyridazinecarboxamide, 3-chloro-6-methyl- (7CI, 8CI) (CA INDEX NAME)

$$\begin{array}{c|c} & & & \text{Cl} \\ & & & \\ \text{Me} & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ &$$

RN 2125-88-4 CAOLD

CN 4-Pyridazinecarboxamide, 3-chloro- (7CI, 8CI, 9CI) (CA INDEX NAME)

L4 ANSWER 2 OF 2 CAOLD COPYRIGHT 2004 ACS on STN

AN CA53:22026b CAOLD

TI 6-chloropyridazines

PA CIBA Ltd.

DT Patent

PATENT NO. KIND DATE

PI GB 807548

IT 7083-49-0 **54108-27-9** 93824-72-7 94011-79-7

101884-08-6 107919-08-4

IT 54108-27-9 101884-08-6

RN 54108-27-9 CAOLD

CN 4-Pyridazinecarboxylic acid, 3-chloro-5,6-diphenyl-, ethyl ester (6CI, 9CI) (CA INDEX NAME)

RN 101884-08-6 CAOLD

CN 4-Pyridazinecarboxylic acid, 3-chloro-5,6-bis(p-chlorophenyl)-, ethyl ester (6CI) (CA INDEX NAME)

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substance identification.

=> s 1341 L3 T₁5 => sort py 15 SORT ENTIRE ANSWER SET? (Y)/N:. PROCESSING COMPLETED FOR L5 41 SORT L5 PY => d 1-20 cbib pi fhistr 'FHISTR' IS NOT A VALID FORMAT FOR FILE 'CAPLUS' The following are valid formats: ABS ----- GI and AB ALL ----- BIB, AB, IND, RE APPS ----- AI, PRAI BIB ----- AN, plus Bibliographic Data and PI table (default) CAN ----- List of CA abstract numbers without answer numbers CBIB ----- AN, plus Compressed Bibliographic Data DALL ---- ALL, delimited (end of each field identified) DMAX ----- MAX, delimited for post-processing FAM ----- AN, PI and PRAI in table, plus Patent Family data FBIB ----- AN, BIB, plus Patent FAM IND ----- Indexing data IPC ----- International Patent Classifications MAX ----- ALL, plus Patent FAM, RE PATS ----- PI, SO SAM ----- CC, SX, TI, ST, IT SCAN ----- CC, SX, TI, ST, IT (random display, no answer numbers; SCAN must be entered on the same line as the DISPLAY, e.g., D SCAN or DISPLAY SCAN) STD ---- BIB, IPC, and NCL IABS ----- ABS, indented with text labels IALL ----- ALL, indented with text labels IBIB ----- BIB, indented with text labels IMAX ----- MAX, indented with text labels ISTD ----- STD, indented with text labels OBIB ----- AN, plus Bibliographic Data (original) OIBIB ----- OBIB, indented with text labels SBIB ----- BIB, no citations SIBIB ----- IBIB, no citations HIT ----- Fields containing hit terms HITIND ----- IC, ICA, ICI, NCL, CC and index field (ST and IT) containing hit terms HITRN ----- HIT RN and its text modification HITSTR ----- HIT RN, its text modification, its CA index name, and

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HITSEQ ----- HIT RN, its text modification, its CA index name, its structure diagram, plus NTE and SEQ fields

FHITSTR ---- First HIT RN, its text modification, its CA index name, and its structure diagram

FHITSEQ ---- First HIT RN, its text modification, its CA index name, its structure diagram, plus NTE and SEQ fields

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L6 ANSWER 1 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN

1959:122326 Document No. 53:122326 Original Reference No. 53:22026c-d Isotope-tagged 5-ethyl-5-(1-methylbutyl)thiobarbituric acid and its sodium salt. Anorova, G. A.; Shishkov, V. P. SU 118557 19590310 (Unavailable). APPLICATION: SU .

PATENT NO. KIND DATE APPLICATION NO. DATE

PI SU 118557 19590310 SU

(preparation of)

RN 54108-27-9 CAPLUS

CN 4-Pyridazinecarboxylic acid, 3-chloro-5,6-diphenyl-, ethyl ester (6CI, 9CI) (CA INDEX NAME)

L6 ANSWER 2 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN

1959:122325 Document No. 53:122325 Original Reference No. 53:22026b-c 6-Chloropyridazines. (C I B A Ltd.). GB 807548 19590114 (Unavailable). APPLICATION: GB.

PATENT NO. KIND DATE APPLICATION NO. DATE

PI GB 807548 19590114 GB

IT 101884-08-6, 4-Pyridazinecarboxylic acid, 3-chloro-5,6-bis(pchlorophenyl)-, ethyl ester

(preparation of)

RN 101884-08-6 CAPLUS

CN 4-Pyridazinecarboxylic acid, 3-chloro-5,6-bis(p-chlorophenyl)-, ethyl ester (6CI) (CA INDEX NAME)

L6 ANSWER 3 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN

1965:36816 Document No. 62:36816 Original Reference No. 62:6480a-f Synthesis of nitrogen-containing heterocycles. XXXII. Syntheses of pyrazolo[3.4-c]pyridazines. Dornow, Alfred; Abele, Werner (Tech. Hochschule, Hannover, Germany). Ber., 97(12), 3349-53 (German) 1964. OTHER SOURCES: CASREACT 62:36816.

IT 1445-53-0, 4-Pyridazinecarboxylic acid, 3-chloro-6-methyl-, ethyl ester

(preparation of)

RN 1445-53-0 CAPLUS

CN 4-Pyridazinecarboxylic acid, 3-chloro-6-methyl-, ethyl ester (7CI, 8CI) (CA INDEX NAME)

L6 ANSWER 4 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN

1969:470623 Document No. 71:70623 Coccidiocidal pyridazin-5-yl-methyl-2-picolinium halides. Ramuz, Henri; Spiegelberg, Hans; Abele, Werner; Dornow, Alfred (Hoffmann-La Roche Inc.). U.S. US 3453277 19690701, 5 pp.

(English). CODEN: USXXAM. APPLICATION: US 1966-539580 19660401.

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE		
		_					
ΡI	US 3453277	Α	19690701	US 1966-539580	19660401		
	BE 679422	Α	19661013	BE 1966-679422	19660413		
	NL 6605001	Α	19661017	NL 1966-5001	19660414		

IT 1445-53-0P

RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)

RN 1445-53-0 CAPLUS

CN 4-Pyridazinecarboxylic acid, 3-chloro-6-methyl-, ethyl ester (7CI, 8CI) (CA INDEX NAME)

L6 ANSWER 5 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN

1972:34229 Document No. 76:34229 Syntheses of N-heterocyclic compounds. II. Pyrimido[4,5-e]-, pyridazino[3,4-e]-, and pyrido[4,3-e]-1,2,3,5-tetrahydro[1,4]oxazepin-5-one. Yurugi, Shojiro; Hieda, Masaru; Fushimi, Tomiyoshi; Tomimoto, Mitsumi (Res. Dev. Div., Takeda Chem. Ind., Ltd., Osaka, Japan). Chemical & Pharmaceutical Bulletin, 19(11), 2354-64 (English) 1971. CODEN: CPBTAL. ISSN: 0009-2363.

IT 34750-70-4P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation and cycloaddn. reaction of, with ethanolamines)

RN 34750-70-4 CAPLUS

CN 4-Pyridazinecarboxylic acid, 3-chloro-6-phenyl-, ethyl ester (9CI) (CA INDEX NAME)

L6 ANSWER 6 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN

1974:552256 Document No. 81:152256 3-[(Substituted)amino]-5,6-diphenyl-4-pyridazinecarboxylic acids. Santilli, Arthur A.; Scotese, Anthony C.; Kim, Dong H. (American Home Products Corp.). U.S. US 3836532 19740917, 2 pp. (English). CODEN: USXXAM. APPLICATION: US 1972-302303 19721030.

PATENT NO. KIND DATE APPLICATION NO. DATE

I US 3836532 A 19740917 US 1972-302303 19721030

IT 54108-27-9P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation and reaction with amines)

RN 54108-27-9 CAPLUS

CN 4-Pyridazinecarboxylic acid, 3-chloro-5,6-diphenyl-, ethyl ester (6CI, 9CI) (CA INDEX NAME)

L6 ANSWER 7 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN

1977:601446 Document No. 87:201446 Studies on heterocyclic compounds. XX. The reaction of 4-cyano-3,6-dichloropyridazine with amines. Yanai, Mitsuji; Takeda, Shigeko; Mitsuoka, Tamao (Fac. Pharm. Sci., Nagasaki Univ., Nagasaki, Japan). Chemical & Pharmaceutical Bulletin, 25(7), 1708-16 (English) 1977. CODEN: CPBTAL. ISSN: 0009-2363. OTHER SOURCES: CASREACT 87:201446.

IT 64613-29-2P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation and dechlorination of)

RN 64613-29-2 CAPLUS

CN 4-Pyridazinecarboxamide, 3-chloro-6-(4-morpholinyl)- (9CI) (CA INDEX NAME)

$$\begin{array}{c|c} & & & \\ & & \\ & & & \\ & & & \\ & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ &$$

L6 ANSWER 8 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN

1979:456936 Document No. 91:56936 Synthesis and pharmacological activity of some 5,6-diphenyl-pyridazines. Nannini, Giuliano; Biasoli, Giovanni; Perrone, Ettore; Forgione, Angelo; Buttinoni, Ada; Ferrari, Mario (Carlo Erba Res. Inst., Milan, 20159, Italy). European Journal of Medicinal Chemistry, 14(1), 53-60 (English) 1979. CODEN: EJMCA5. ISSN: 0009-4374. OTHER SOURCES: CASREACT 91:56936.

IT 70769-18-5P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation and reactions of)

RN 70769-18-5 CAPLUS

CN 4-Pyridazinecarboxylic acid, 3-chloro-5,6-bis(4-methoxyphenyl)-, ethyl ester (9CI) (CA INDEX NAME)

L6 ANSWER 9 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN

1986:626078 Document No. 105:226078 Benzoylurea derivatives having antitumor activity. Brouwer, Marius S.; Van Hes, Roelof (Duphar International Research B. V., Neth.). Eur. Pat. Appl. EP 193249 A2 19860903, 31 pp. DESIGNATED STATES: R: AT, BE, CH, DE, FR, GB, IT, LI, LU, NL, SE. (English). CODEN: EPXXDW. APPLICATION: EP 1986-200300 19860227. PRIORITY: NL 1985-572 19850301.

PAT	PATENT NO.			DATE	APPLICATION NO.	DATE		
PI EP	193249		A2	19860903	EP 1986-200300	19860227		
EP	193249		A3	19880316				
	R: AT,	BE, C	CH, DE	, FR, GB, I	T, LI, LU, NL, SE			
DK	8600881		Α	19860902	DK 1986-881	19860226		
AU	8654108		A1	19860904	AU 1986-54108	19860226		
AU	601145		B2	19900906				
ZA	8601446		Α	19861029	ZA 1986-1446	19860226		
ES	552432		A1	19880301	ES 1986-552432	19860226		
JP	61218569		A2	19860929	JP 1986-42838	19860301		

IT 105390-24-7P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (preparation of, as antitumor agent)

RN 105390-24-7 CAPLUS

CN 4-Pyridazinecarboxamide, 3-chloro-N-[[[4-(trifluoromethyl)phenyl]amino]carbonyl]- (9CI) (CA INDEX NAME)

L6 ANSWER 10 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN 1986:478949 Document No. 105:78949 3-[(Methyl or dimethyl)amino]-6-(pyridinyl)pyridazines and their cardiotonic use. Lesher, George Y.;

Dickinson, William B. (Sterling Drug Inc., USA). U.S. US 4590194 A 19860520, 7 pp. (English). CODEN: USXXAM. APPLICATION: US 1981-308413 19811005.

IT 103556-15-6P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation and reaction of)

RN 103556-15-6 CAPLUS

CN 4-Pyridazinecarboxamide, 3-chloro-6-(4-pyridinyl)- (9CI) (CA INDEX NAME)

L6 ANSWER 11 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN

1988:75327 Document No. 108:75327 Pyridazine chemistry. XXXIV. Synthesis of novel diaza analogs of acridone, xanthone and thioxanthone from 3,4-disubstituted pyridazines. Heinisch, Gottfried; Lassnigg, Doris (Inst. Pharm. Chem., Univ. Wien, Vienna, A-1090, Austria). Archiv der Pharmazie (Weinheim, Germany), 320(12), 1222-6 (German) 1987. CODEN: ARPMAS. ISSN: 0365-6233. OTHER SOURCES: CASREACT 108:75327.

IT 112584-71-1P

RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)

RN 112584-71-1 CAPLUS

CN 4-Pyridazinecarboxamide, 3-phenoxy- (9CI) (CA INDEX NAME)

$$\begin{array}{c|c} N & \text{OPh} \\ \hline \\ C-NH_2 \\ \hline \\ O \end{array}$$

L6 ANSWER 12 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN

1988:56021 Document No. 108:56021 Syntheses of 4-aroyl- and 4-[disubstituted (hydroxymethyl)]pyridazin-3(2H)-ones and their 3-mercapto derivatives: synthesis of 1,3-dithiano[4,5-c]pyridazine as a new ring system. Shams, Nabil A.; Enayat, Ebtisam I.; Shalaby, Alyaa A.; El-Abbady, Samia A. (Fac. Sci., Ain Shams Univ., Cairo, Egypt). Journal fuer Praktische Chemie (Leipzig), 329(2), 271-6 (English) 1987. CODEN: JPCEAO. ISSN: 0021-8383. OTHER SOURCES: CASREACT 108:56021.

IT 112450-33-6

RL: RCT (Reactant); RACT (Reactant or reagent) (Grignard reactions of, with alkyl halides)

RN 112450-33-6 CAPLUS

4-Pyridazinecarboxylic acid, 2,3-dihydro-5,6-diphenyl-3-thioxo-, ethyl CNester (9CI) (CA INDEX NAME)

ANSWER 13 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN

1988:6047 Document No. 108:6047 Preparation of (heterocyclylpiperazinyl)alka nols as antitussives. Giani, Roberto P.; Malandrino, Salvatore; Tonon, Giancarlo (Dompe Farmaceutici S.p.A., Italy). Eur. Pat. Appl. EP 230402 A2 19870729, 24 pp. DESIGNATED STATES: R: AT, BE, CH, DE, ES, FR, GB, GR, IT, LI, LU, NL, SE. (English). CODEN: EPXXDW. APPLICATION: EP 1987-830012 19870116. PRIORITY: IT 1986-19121 19860120.

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡI	EP 230402	A2	19870729	EP 1987-830012	19870116
	EP 230402	A3	19890222		
	EP 230402	В1	19930421		
	R: AT, BE,	CH, DE	, ES, FR, GB,	GR, IT, LI, LU, NL	, SE
	US 4849422	A	19890718	US 1987-4028	19870116
	AT 88467	E	19930515	AT 1987-830012	19870116
	ES 2056836	Т3	19941016	ES 1987-830012	19870116
	JP 62169768	A2	19870725	JP 1987-8371	19870119

IT1445-54-1

RL: RCT (Reactant); RACT (Reactant or reagent) (amination of, by (dihydroxypropyl)piperazine)

1445-54-1 CAPLUS RN

4-Pyridazinecarboxylic acid, 3-chloro-, ethyl ester (7CI, 8CI, 9CI) (CA CN INDEX NAME)

112450-33-6 CAPLUS

- L6 ANSWER 14 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN
- 1990:478301 Document No. 113:78301 Action of acrylonitrile on 4-cyano- and 4-(carbethoxy)pyridazin-3(2H)-ones and their 3-mercapto derivatives. A novel synthesis of pyrano- and thiinopyridazines. Shams, Nabil A.; Kaddah, Ahmed M.; Enayat, Ebtisam I.; Shalaby, Alyaa A.; Youssef, Ali M. (Fac. Sci., Ain Shams Univ., Cairo, Egypt). Journal fuer Praktische Chemie (Leipzig), 331(5), 757-62 (English) 1989. CODEN: JPCEAO. ISSN: 0021-8383. OTHER SOURCES: CASREACT 113:78301.
- IT 112450-33-6

RN

RL: RCT (Reactant); RACT (Reactant or reagent) (addition and cycloaddn. reactions with acrylonitrile and hydrolysis of)

CN 4-Pyridazinecarboxylic acid, 2,3-dihydro-5,6-diphenyl-3-thioxo-, ethyl ester (9CI) (CA INDEX NAME)

- L6 ANSWER 15 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN
- 1989:114776 Document No. 110:114776 3-Aminopyridazine derivatives with atypical antidepressant, serotonergic and dopaminergic activities. Wermuth, Camille Georges; Schlewer, Gilbert; Bourguignon, Jean Jacques; Maghioros, Georges; Bouchet, Marie Jeanne; Moire, Claudine; Kan, Jean Paul; Worms, Paul; Biziere, Kathleen (Dep. Pharmacochim. Mol., Univ. Louis Pasteur, Strasbourg, 67084, Fr.). Journal of Medicinal Chemistry, 32(3), 528-37 (English) 1989. CODEN: JMCMAR. ISSN: 0022-2623. OTHER SOURCES: CASREACT 110:114776.
- IT 34750-70-4P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation and amination and amidation of)

- RN 34750-70-4 CAPLUS
- CN 4-Pyridazinecarboxylic acid, 3-chloro-6-phenyl-, ethyl ester (9CI) (CA INDEX NAME)

L6 ANSWER 16 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN

1990:497555 Document No. 113:97555 Pyridazine analogs of biologically active compounds. VI. Pyrido[2,3-c]pyridazines structurally related to nalidixic acid. Boamah, Philip Y.; Haider, Norbert; Heinisch, Gottfried

(Inst. Pharm. Chem., Univ. Vienna, Vienna, A-1090, Austria). Archiv der Pharmazie (Weinheim, Germany), 323(4), 207-10 (English) 1990. CODEN: ARPMAS. ISSN: 0365-6233. OTHER SOURCES: CASREACT 113:97555.

IT 1445-54-1

RL: RCT (Reactant); RACT (Reactant or reagent) (amination of, with propylaminopropionate)

- RN 1445-54-1 CAPLUS
- CN 4-Pyridazinecarboxylic acid, 3-chloro-, ethyl ester (7CI, 8CI, 9CI) (CA INDEX NAME)

- L6 ANSWER 17 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN
- 1990:478302 Document No. 113:78302 Reactions of ethyl monohaloacetate and hydrazine hydrate with 4-cyano and 4-carbethoxypyridazin-3(2H)-ones and their 3-mercapto derivatives. Syntheses of furo, thieno and pyrazolo pyridazines. Shalaby, Alyaa A. (Fac. Sci., Ain Shams Univ., Cairo, Egypt). Journal fuer Praktische Chemie (Leipzig), 332(1), 104-8 (English) 1990. CODEN: JPCEAO. ISSN: 0021-8383. OTHER SOURCES: CASREACT 113:78302.
- IT 112450-33-6

RL: RCT (Reactant); RACT (Reactant or reagent) (cyclocondensation of, with haloacetate and hydrazine)

- RN 112450-33-6 CAPLUS
- CN 4-Pyridazinecarboxylic acid, 2,3-dihydro-5,6-diphenyl-3-thioxo-, ethyl ester (9CI) (CA INDEX NAME)

L6 ANSWER 18 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN

1991:471518 Document No. 115:71518 Pyridazines. LIV. The synthesis of pyridazine-fused S-heterocycles: thieno[2,3-c]pyridazine, pyrimido[4',5':4,5]thieno[2,3-c]pyridazine, and pyridazino[3,4-b][1,4]benzothiazine. Czech, Karin; Haider, Norbert; Heinisch, Gottfried (Inst. Pharm. Chem., Univ. Vienna, Vienna, A-1090, Austria). Monatshefte fuer Chemie, 122(5), 413-18 (English) 1991. CODEN: MOCMB7. ISSN: 0026-9247. OTHER SOURCES: CASREACT 115:71518.

IT 1445-54-1

RL: RCT (Reactant); RACT (Reactant or reagent)
 (cyclization of, with aminothiophenol)

RN 1445-54-1 CAPLUS

CN 4-Pyridazinecarboxylic acid, 3-chloro-, ethyl ester (7CI, 8CI, 9CI) (CA INDEX NAME)

L6 ANSWER 19 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN

1995:9882 Document No. 122:31861 Synthesis of 4-cyano, 4-carboethoxy and 4-carboxy pyridazine amino acid derivatives of expected biological activity. Ibrahim, T. M.; Donia, S. G.; Magdel-Din, A. A. (Fac. Sci., Al-Azhar Univ., Nasr City, Egypt). Al-Azhar Bulletin of Science, 3(2), 423-32 (English) 1992. CODEN: ABSCE7. ISSN: 1110-2535.

IT 159580-20-8P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)

(preparation and bactericidal and fungicidal activity of)

RN 159580-20-8 CAPLUS

CN 4-Pyridazinecarboxylic acid, 3-[[3-mercapto-2-[[(4-methylphenyl)sulfonyl]amino]-1-oxopropyl]thio]-5,6-diphenyl-, (R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

L6 ANSWER 20 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN

1992:426497 Document No. 117:26497 Enzymic hydrolysis of heterocyclic nitriles. Klempier, Norbert; De Raadt, Anna; Griengl, Herfried; Heinisch, Gottfried (Inst. Org. Chem., Graz Univ. Technol., Graz, A-8010, Austria). Journal of Heterocyclic Chemistry, 29(1), 93-5 (English) 1992. CODEN: JHTCAD. ISSN: 0022-152X. OTHER SOURCES: CASREACT 117:26497.

IT 2125-88-4P

RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)

RN 2125-88-4 CAPLUS

CN 4-Pyridazinecarboxamide, 3-chloro- (7CI, 8CI, 9CI) (CA INDEX NAME)

=> d 21-30 cbib pi fhitstr

L6 ANSWER 21 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN

1994:77242 Document No. 120:77242 A new two-step procedure for functionalized 3(2H)-pyridazinones through homolytic substitution of 3-chloropyridazines. Dal Piaz, Vittorio; Giovannoni, Maria Paola; Ciciani, Giovanna (Dip. Sci. Farm., Univ. Florence, Florence, I-50121, Italy). Tetrahedron Letters, 34(24), 3903-6 (English) 1993. CODEN: TELEAY. ISSN: 0040-4039. OTHER SOURCES: CASREACT 120:77242.

IT 152343-20-9P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent) (preparation and hydrolysis of)

RN 152343-20-9 CAPLUS

CN 4,5-Pyridazinedicarboxylic acid, 3-chloro-6-methyl-, diethyl ester (9CI) (CA INDEX NAME)

L6 ANSWER 22 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN 1993:625966 Document No. 119:225966 Preparative and biological activity of aryl substituted nitrogen containing heterocycles. Linz, Guenter; Pieper,

Helmut; Himmelsbach, Frank; Austel, Volkhard; Mueller, Thomas; Weisenberger, Johannes; Seewaldt-Becker, Elke (Thomae, Dr. Karl, G.m.b.H., Germany). Eur. Pat. Appl. EP 537696 A1 19930421, 47 pp. DESIGNATED STATES: R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, NL, PT, SE. (German). CODEN: EPXXDW. APPLICATION: EP 1992-117507 19921014. PRIORITY: DE 1991-4134467 19911018.

	PATENT NO.	ATENT NO. KIND DATE		APPLICATION NO.	DATE
			_		
ΡI	EP 537696	A1	19930421	EP 1992-117507	19921014
	R: AT, BE,	CH, DE	, DK, ES, FR,	GB, GR, IE, IT, LI	, LU, NL, PT, SE
,	DE 4134467	A 1	19930422	DE 1991-4134467	19911018
	US 5418233	Α	19950523	US 1992-961135	19921014
	CA 2080748	AA	19930419	CA 1992-2080748	19921016
	NO 9204027	Α	19930419	NO 1992-4027	19921016
	AU 9227062	A1	19930422	AU 1992-27062	19921016
	AU 662930	B2	19950921		
	HU 62272	A2	19930428	HU 1992-3264	19921016
	JP 05221992	A2	19930831	JP 1992-277578	19921016
	ZA 9207994	Α	19940418	ZA 1992-7994	19921016
	US 5563268	Α	19961008	US 1995-375084	19950119

IT 150594-29-9P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation and reaction of, in preparation of thromboxane formation inhibitor)

RN 150594-29-9 CAPLUS

CN 4-Pyridazinecarboxylic acid, 3-chloro-6-(4-cyanophenyl)-, ethyl ester (9CI) (CA INDEX NAME)

L6 ANSWER 23 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN 1993:213015 Document No. 118:213015 Heterocyclic β-enamino esters. 54. A new, general approach for the synthesis of heteroannulated 3,1-oxazin-4-ones. Wamhoff, Heinrich; Herrmann, Stefan; Stoelben, Stephan; Nieger, Martin (Inst. Org. Chem. Biochem., Univ. Bonn, Bonn, D-5300, Germany). Tetrahedron, 49(3), 581-94 (English) 1993. CODEN: TETRAB. ISSN: 0040-4020. OTHER SOURCES: CASREACT 118:213015.

IT 54108-27-9

RL: RCT (Reactant); RACT (Reactant or reagent) (sequential azidation and Staudinger reaction with triphenylphosphine) 54108-27-9 CAPLUS

CN 4-Pyridazinecarboxylic acid, 3-chloro-5,6-diphenyl-, ethyl ester (6CI, 9CI) (CA INDEX NAME)

L6 ANSWER 24 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN

1995:432038 Document No. 123:112646 Synthesis of 4-cyano-, 4-carbethoxy- and 4-carboxy pyridazine amino acid derivatives of expected biological activity. Ibrahim, T. M.; Donia, S. G.; Magdel-Din, A. A. (Faculty Science, Al-Azhar and Benha Universities, Cairo, Egypt). Egyptian Journal of Chemistry, 37(3), 273-81 (English) 1994. CODEN: EGJCA3. ISSN: 0367-0422. Publisher: National Information and Documentation Centre.

IT 159579-98-3P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); RCT (Reactant); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent) (preparation and antimicrobial activity of cyano-, carbethoxy-, and

carboxypyridazine amino acid esters and thioesters)

RN 159579-98-3 CAPLUS

CN 4-Pyridazinecarboxylic acid, 3-[[(1,3-dihydro-1,3-dioxo-2H-isoindol-2-yl)acetyl]thio]-5,6-diphenyl-, ethyl ester (9CI) (CA INDEX NAME)

L6 ANSWER 25 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN 1995:382788 Document No. 122:160663 Preparation of 6-(hetero)ary1-3-chloroyridazine-4-carboxamides and analogs as interleukin-1β protease inhibitors. Dolle, Roland E.; Hoyer, Denton W.; Schmidt, Stanley J.; Ross, Tina M.; Rinker, James M.; Ator, Mark A. (Sterling Winthrop Inc., USA). Eur. Pat. Appl. EP 628550 A2 19941214, 18 pp. DESIGNATED STATES: R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE. (English). CODEN: EPXXDW. APPLICATION: EP 1994-201597 19940604. PRIORITY: US 1993-73914 19930608.

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡI	EP 628550	A2	19941214	EP 1994-201597	19940604
	EP 628550	A3	19950118		
	EP 628550	B1	19980225		
	R: AT, BE,	CH, DE	, DK, ES, FR,	GB, GR, IE, IT, LI	, LU, MC, NL, PT, SE
	AT 163412	E	19980315	AT 1994-201597	19940604
	ES 2114654	Т3	19980601	ES 1994-201597	19940604

2125359	AA	19941209			19940607
9464556	A1	19941215	AU	1994-64556	19940607
693786	B2	19980709		¥	
07069894	A2	19950314	JΡ	1994-125091	19940607
9402693	Α	19941209	FI	1994-2693	19940608
9402133	Α	19941209	NO	1994-2133	19940608
67413	A2	19950428	HU	1994-1714	19940608
6121266	Α	20000919	US	1995-444984	19950519
6624166	В1	20030923	US	1996-625680	19960403
	9464556 593786 57069894 9402693 9402133 57413 5121266	0464556 A1 593786 B2 07069894 A2 0402693 A 0402133 A 07413 A2 0121266 A	0464556 A1 19941215 593786 B2 19980709 07069894 A2 19950314 0402693 A 19941209 0402133 A 19941209 07413 A2 19950428 0121266 A 20000919	9464556 A1 19941215 AU 593786 B2 19980709 57069894 A2 19950314 JP 6402693 A 19941209 FI 6402133 A 19941209 NO 57413 A2 19950428 HU 5121266 A 20000919 US	9464556 A1 19941215 AU 1994-64556 693786 B2 19980709 97069894 A2 19950314 JP 1994-125091 9402693 A 19941209 FI 1994-2693 9402133 A 19941209 NO 1994-2133 67413 A2 19950428 HU 1994-1714 6121266 A 20000919 US 1995-444984

IT 103556-15-6P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of 6-(hetero)aryl-3-chloropyridazine-4-carboxamides and analogs as interleukin-1 β protease inhibitors)

RN 103556-15-6 CAPLUS

CN 4-Pyridazinecarboxamide, 3-chloro-6-(4-pyridinyl)- (9CI) (CA INDEX NAME)

L6 ANSWER 26 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN

1997:27720 Document No. 126:144238 Synthesis and reactivity of some pyridazine derivatives. Kamal El-Dean, A. M.; Radwan, Sh. M.; Abdel Moneam, M. I. (Fac. Sci., Assiut Univ., Assiut, 71516, Egypt). Afinidad, 53(466), 387-391 (English) 1996. CODEN: AFINAE. ISSN: 0001-9704. Publisher: Asociacion de Quimicos del Instituto Quimico de Sarria.

IT 54108-27-9P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of pyridazine derivs.)

RN 54108-27-9 CAPLUS

CN 4-Pyridazinecarboxylic acid, 3-chloro-5,6-diphenyl-, ethyl ester (6CI, 9CI) (CA INDEX NAME)

L6 ANSWER 27 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN 1997:558055 Document No. 127:262667 Preparation of pyrazolo[1,5-a]pyridine

derivatives as adenosine antagonists and their pharmaceutical uses. Kuroda, Satoshi; Itani, Hiromichi; Akabane, Atsushi (Fujisawa Pharmaceutical Co., Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 09216883 A2 19970819 Heisei, 17 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1996-24146 19960209.

PATENT NO. KIND DATE APPLICATION NO. DATE

----PI JP 09216883 A2 19970819 JP 1996-24146 19960209

IT 195827-20-4P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of pyrazolo[1,5-a]pyridine derivs. as adenosine antagonists and their pharmaceutical uses)

RN 195827-20-4 CAPLUS

CN 4-Pyridazinecarboxylic acid, 3-chloro-6-(2-phenylpyrazolo[1,5-a]pyridin-3-yl)-, ethyl ester (9CI) (CA INDEX NAME)

L6 ANSWER 28 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN

1997:433272 Document No. 127:121687 Pyridazine derivatives. Synthesis and reactions of some new indeno[1,2-d]pyridazine derivatives. Radwan, S. M.; El-Dean, A. M. Kamal (Faculty Science, Assiut University, Assiut, 71516, Egypt). Pharmazie, 52(6), 483-485 (English) 1997. CODEN: PHARAT. ISSN: 0031-7144. OTHER SOURCES: CASREACT 127:121687. Publisher: Govi-Verlag Pharmazeutischer Verlag.

IT **54108-27-9**

RL: RCT (Reactant); RACT (Reactant or reagent)
 (preparation of indenopyridazines)

RN 54108-27-9 CAPLUS

CN 4-Pyridazinecarboxylic acid, 3-chloro-5,6-diphenyl-, ethyl ester (6CI, 9CI) (CA INDEX NAME)

L6 ANSWER 29 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN

1997:407617 Document No. 127:121686 Pyridazine N-oxides. II. Synthesis and in vitro antimicrobial evaluation of 3-chloro-4-carbamoyl-5-aryl-6-methylpyridazine N-oxides. Gavini, Elisabetta; Juliano, Claudia; Mule, Antonio; Pirisino, Gerolamo; Pinna, Gerard A. (Dip. Scienze Farmaco, Muroni, Sassari, 07100, Italy). Farmaco, 52(1), 67-69 (English) 1997. CODEN: FRMCE8. ISSN: 0014-827X. Publisher: Societa Chimica Italiana.

IT 192634-55-2P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)

(preparation and antimicrobial evaluation of)

RN 192634-55-2 CAPLUS

CN 4-Pyridazinecarboxamide, 3-chloro-6-methyl-N,5-diphenyl-, 1-oxide (9CI) (CA INDEX NAME)

L6 ANSWER 30 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN

1997:330887 Document No. 127:65731 3-Chloro-4-carboxamido-6-arylpyridazines as a non-peptide class of interleukin-1β converting enzyme inhibitor. Dolle, Roland E.; Hoyer, Denton; Rinker, James M.; Ross, Tina Morgan; Schmidt, Stanley J.; Helaszek, Carla; Ator, Mark A. (Sanofi Winthrop Inc., Malvern, PA, 19426, USA). Bioorganic & Medicinal Chemistry Letters, 7(8), 1003-1006 (English) 1997. CODEN: BMCLE8. ISSN: 0960-894X. Publisher: Elsevier.

IT 161228-78-0

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study)

(preparation of (aryl)chloropyridazinecarboxamides as interleukin 1β -converting enzyme inhibitors)

RN 161228-78-0 CAPLUS

CN 4-Pyridazinecarboxylic acid, 3-chloro-6-(4-pyridinyl)-, ethyl ester (9CI) (CA INDEX NAME)

=> d 11 cbib pi hitstr

L6 ANSWER 11 OF 41 CAPLUS COPYRIGHT 2004 ACS on STN 1988:75327 Document No. 108:75327 Pyridazine chemistry. XXXIV. Synthesis of novel diaza analogs of acridone, xanthone and thioxanthone from 3,4-disubstituted pyridazines. Heinisch, Gottfried; Lassnigg, Doris (Inst. Pharm. Chem., Univ. Wien, Vienna, A-1090, Austria). Archiv der Pharmazie (Weinheim, Germany), 320(12), 1222-6 (German) 1987. CODEN: ARPMAS. ISSN: 0365-6233. OTHER SOURCES: CASREACT 108:75327.

RN 112584-71-1 CAPLUS

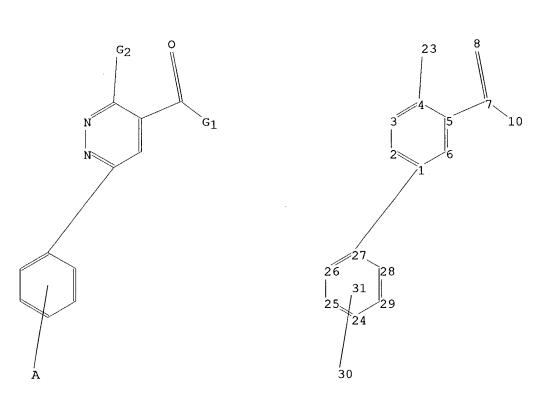
CN 4-Pyridazinecarboxamide, 3-phenoxy- (9CI) (CA INDEX NAME)

$$\begin{array}{c|c} N & \text{OPh} \\ \hline \\ C-NH_2 \\ \hline \\ O \end{array}$$

RN 112584-72-2 CAPLUS CN 4-Pyridazinecarboxylic acid, 3-phenoxy- (9CI) (CA INDEX NAME)

=> Uploading C:\Program Files\Stnexp\Queries\10668475.str





chain nodes : 7 8 10 17 18 19 23 30 ring nodes : 1 2 3 4 5 6 11 12 13 14 15 16 24 25 26 27 28 29 chain bonds : 1-27 4-23 5-7 7-8 7-10 14-17 18-19 ring bonds : $1 - 2 \quad 1 - 6 \quad 2 - 3 \quad 3 - 4 \quad 4 - 5 \quad 5 - 6 \quad 11 - 12 \quad 11 - 16 \quad 12 - 13 \quad 13 - 14 \quad 14 - 15 \quad 15 - 16 \quad 24 - 25$ 24-29 25-26 26-27 27-28 28-29 exact/norm bonds : 4-23 7-8 7-10 14-17 18-19 exact bonds : 1-27 5-7 normalized bonds : $1-2 \quad 1-6 \quad 2-3 \quad 3-4 \quad 4-5 \quad 5-6 \quad 11-12 \quad 11-16 \quad 12-13 \quad 13-14 \quad 14-15 \quad 15-16 \quad 24-25$ 24-29 25-26 26-27 27-28 28-29

G1:0,N

G2:X,S,P,[*1],[*2]

Match level:

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:CLASS 8:CLASS 10:CLASS 11:Atom 12:Atom 13:Atom 14:Atom 15:Atom 16:Atom 17:CLASS 18:CLASS 19:CLASS 23:CLASS 24:CLASS 25:Atom 26:Atom 27:Atom 28:Atom 29:Atom 30:CLASS 31:CLASS

L7 STRUCTURE UPLOADED

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=> s 17 subset = 13
ENTER SUBSET SEARCH SCOPE - SAMPLE, FULL, RANGE, OR (END):sample
SAMPLE SUBSET SEARCH INITIATED 13:12:26 FILE 'REGISTRY'
SAMPLE SUBSET SCREEN SEARCH COMPLETED - 2 TO ITERATE

100.0% PROCESSED 2 ITERATIONS 1 ANSWERS SEARCH TIME: 00.00.01

PROJECTIONS (WITHIN SPECIFIED SUBSET): ONLINE **COMPLETE**
PROJECTED ITERATIONS (WITHIN SPECIFIED SUBSET): 2 TO 124
PROJECTED ANSWERS (WITHIN SPECIFIED SUBSET): 1 TO 80

L8 1 SEA SUB=L3 SSS SAM L7

=> d scan

L8 1 ANSWERS REGISTRY COPYRIGHT 2004 ACS on STN

4-Pyridazinecarboxamide, 3-chloro-6-(4-cyanophenyl)- (9CI) IN C12 H7 Cl N4 O MF

$$\begin{array}{c|c} C1 & N & N & CN \\ H_2N-C & \parallel & & \\ O & & & \end{array}$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

ALL ANSWERS HAVE BEEN SCANNED

=> s 17 subset = 13 full FULL SUBSET SEARCH INITIATED 13:12:46 FILE 'REGISTRY' FULL SUBSET SCREEN SEARCH COMPLETED - 88 TO ITERATE

88 ITERATIONS 100.0% PROCESSED SEARCH TIME: 00.00.01

23 ANSWERS

23 SEA SUB=L3 SSS FUL L7

=> d 1-23 ide cbib pi

ANSWER 1 OF 23 REGISTRY COPYRIGHT 2004 ACS on STN L9

405224-31-9 REGISTRY RN

4-Pyridazinecarboxamide, 3-chloro-6-(2,3-difluorophenyl)- (9CI) (CA INDEX CN NAME)

FS 3D CONCORD

MF C11 H6 C1 F2 N3 O

SR

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 2 REFERENCES IN FILE CA (1907 TO DATE)
- 2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

```
REFERENCE 1: 139:292257 Preparation of pyrazolopyridazines for the
     inhibition of GSK-3. Witherington, Jason; Ward, Robert William (Glaxo
     Group Limited, UK). PCT Int. Appl. WO 2003080616 A1 20031002, 72 pp.
     DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ,
     CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE,
     GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS,
     LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO,
     RU, SC, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ,
     VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE,
     BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT,
     LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English). CODEN:
     PIXXD2. APPLICATION: WO 2003-EP3171 20030320. PRIORITY: GB 2002-6723
     20020321.
                                                   APPLICATION NO. DATE
     PATENT NO.
                          KIND DATE
                                 _____
                                 20031002
                                                   WO 2003-EP3171
                                                                        20030320
     WO 2003080616
                           A1
PΙ
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               CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
               GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
               LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
               PL, PT, RO, RU, SC, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT,
               TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ,
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          RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG,
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               GW, ML, MR, NE, SN, TD, TG
REFERENCE 2: 136:263167 Preparation of pyrazolopyridines and
     pyrazolopyridazines as antidiabetics. Garland, Stephen; Haigh, David;
      Hickey, Deirde Mary Bernadette; Liddle, John; Smith, David Glynn;
      Witherington, Jason; Ward, Robert William (Smithkline Beecham P.L.C., UK).
      PCT Int. Appl. WO 2002024694 A1 20020328, 64 pp. DESIGNATED STATES: W:
     PCT Int. Appl. Wo 2002024694 AI 20020328, 64 pp. DESIGNATED STATES: W:
AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR,
CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID,
IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD,
MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK,
SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG,
KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK,
ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD,
               (English). CODEN: PIXXD2. APPLICATION: WO 2001-GB4186 20010919.
      PRIORITY: GB 2000-23358 20000922; GB 2000-23361 20000922; GB 2001-7391
      20010323; GB 2001-10671 20010501; GB 2001-12799 20010525.
                          KIND DATE
                                                   APPLICATION NO. DATE
      PATENT NO.
                                                                        20010919
                                 20020328
                                                   WO 2001-GB4186
      WO 2002024694
                           A1
               AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
               CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
               GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
               LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL,
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           RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
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                BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
                                                  AU 2001-87898
                                                                        20010919
                           Α5
                                 20020402
      AU 2001087898
                                 20030618
                                                   EP 2001-967523
                                                                        20010919
      EP 1319001
                           Α1
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R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
US 2004019052 A1 20040129 US 2003-381292 20030814

L9 ANSWER 2 OF 23 REGISTRY COPYRIGHT 2004 ACS on STN

RN 343765-74-2 REGISTRY

CN 4-Pyridazinecarboxylic acid, 3-chloro-5,6-bis(4-methoxyphenyl)- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C19 H15 Cl N2 O4

SR Reaction Database

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L9 ANSWER 3 OF 23 REGISTRY COPYRIGHT 2004 ACS on STN

RN 220287-28-5 REGISTRY

CN 4-Pyridazinecarboxylic acid, 3-chloro-6-(4-chlorophenyl)-, ethyl ester (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C13 H10 C12 N2 O2

SR CA

LC STN Files: CA, CAPLUS

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1: 130:153626 Synthesis and pharmacological screening of

pyrazolo[3,4-c]pyridazine derivatives. Zabska, Ryszarda; Kolodziejczyk, Alicja; Sieklucka-Dziuba, Maria; Morawska, Dorota; Kleinrok, Zdzislaw (Department of Chemistry of Drugs, Wroclaw University of Medicine, Wroclaw, 50-137, Pol.). Acta Poloniae Pharmaceutica, 55(4), 305-310 (English) 1998. CODEN: APPHAX. ISSN: 0001-6837. Publisher: Polish Pharmaceutical Society.

L9 ANSWER 4 OF 23 REGISTRY COPYRIGHT 2004 ACS on STN

RN 191411-00-4 REGISTRY

CN 4-Pyridazinecarboxamide, 3-chloro-6-(4-cyanophenyl)- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C12 H7 C1 N4 O

SR CA

LC STN Files: CA, CAPLUS

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1: 127:65731 3-Chloro-4-carboxamido-6-arylpyridazines as a
 non-peptide class of interleukin-1β converting enzyme inhibitor.
 Dolle, Roland E.; Hoyer, Denton; Rinker, James M.; Ross, Tina Morgan;
 Schmidt, Stanley J.; Helaszek, Carla; Ator, Mark A. (Sanofi Winthrop Inc.,
 Malvern, PA, 19426, USA). Bioorganic & Medicinal Chemistry Letters, 7(8),
 1003-1006 (English) 1997. CODEN: BMCLE8. ISSN: 0960-894X. Publisher:
 Elsevier.

L9 ANSWER 5 OF 23 REGISTRY COPYRIGHT 2004 ACS on STN

RN 191410-99-8 REGISTRY

CN 4-Pyridazinecarboxamide, 3-chloro-6-(4-chlorophenyl)- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C11 H7 C12 N3 O

SR CA

LC STN Files: CA, CAPLUS

$$\begin{array}{c|c} C1 & N & N & C1 \\ H_2N-C & \parallel & & \\ O & & & \end{array}$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 1 REFERENCES IN FILE CA (1907 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- REFERENCE 1: 127:65731 3-Chloro-4-carboxamido-6-arylpyridazines as a non-peptide class of interleukin-1β converting enzyme inhibitor.

 Dolle, Roland E.; Hoyer, Denton; Rinker, James M.; Ross, Tina Morgan; Schmidt, Stanley J.; Helaszek, Carla; Ator, Mark A. (Sanofi Winthrop Inc., Malvern, PA, 19426, USA). Bioorganic & Medicinal Chemistry Letters, 7(8), 1003-1006 (English) 1997. CODEN: BMCLE8. ISSN: 0960-894X. Publisher: Elsevier.
- L9 ANSWER 6 OF 23 REGISTRY COPYRIGHT 2004 ACS on STN
- RN 191410-94-3 REGISTRY
- FS 3D CONCORD
- MF C17 H13 C1 N4 O3 S
- SR CA
- LC STN Files: CA, CAPLUS

$$\begin{array}{c|c}
C1 & N & O \\
NH-S-Ph \\
0 & O
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- **PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT**
 - 1 REFERENCES IN FILE CA (1907 TO DATE)
 - 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- REFERENCE 1: 127:65731 3-Chloro-4-carboxamido-6-arylpyridazines as a non-peptide class of interleukin-1β converting enzyme inhibitor. Dolle, Roland E.; Hoyer, Denton; Rinker, James M.; Ross, Tina Morgan; Schmidt, Stanley J.; Helaszek, Carla; Ator, Mark A. (Sanofi Winthrop Inc., Malvern, PA, 19426, USA). Bioorganic & Medicinal Chemistry Letters, 7(8), 1003-1006 (English) 1997. CODEN: BMCLE8. ISSN: 0960-894X. Publisher: Elsevier.
- L9 ANSWER 7 OF 23 REGISTRY COPYRIGHT 2004 ACS on STN
- RN 161228-97-3 REGISTRY
- CN 4-Pyridazinecarboxamide, 3-chloro-6-(3-chloro-4-fluorophenyl)- (9CI) (CA INDEX NAME)
- FS 3D CONCORD
- MF C11 H6 C12 F N3 O
- SR CA
- LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

$$\begin{array}{c|c}
N & & \\
N & & \\
C &$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 2 REFERENCES IN FILE CA (1907 TO DATE)
- 2 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- REFERENCE 1: 127:65731 3-Chloro-4-carboxamido-6-arylpyridazines as a
 non-peptide class of interleukin-1β converting enzyme inhibitor.
 Dolle, Roland E.; Hoyer, Denton; Rinker, James M.; Ross, Tina Morgan;
 Schmidt, Stanley J.; Helaszek, Carla; Ator, Mark A. (Sanofi Winthrop Inc.,
 Malvern, PA, 19426, USA). Bioorganic & Medicinal Chemistry Letters, 7(8),
 1003-1006 (English) 1997. CODEN: BMCLE8. ISSN: 0960-894X. Publisher:
 Elsevier.
- REFERENCE 2: 122:160663 Preparation of 6-(hetero)aryl-3-chloroyridazine-4-carboxamides and analogs as interleukin-1β protease inhibitors.

 Dolle, Roland E.; Hoyer, Denton W.; Schmidt, Stanley J.; Ross, Tina M.; Rinker, James M.; Ator, Mark A. (Sterling Winthrop Inc., USA). Eur. Pat. Appl. EP 628550 A2 19941214, 18 pp. DESIGNATED STATES: R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE. (English). CODEN: EPXXDW. APPLICATION: EP 1994-201597 19940604. PRIORITY: US 1993-73914 19930608.

	PATENT NO.		PATENT NO.			KIND	DATE	APPLICATION NO.			•	DATE				
ΡΊ	EP	628550		A2	19941214		EP	1994	1-20	 1597	_	1994	0604			
	EP	628550		A 3	19950118											
	EΡ	628550		В1	19980225											
		R: AT,	BE,	CH, D	E, DK, ES,	FR,	GB, G	GR, I	Έ, Ξ	ΙΤ,	LI,	LU,	MC,	NL,	PT,	SE
	AT	163412		E	19980315		AT	1994	1-20	1597		1994	0604			
	ES	2114654		Т3	19980601		ES	1994	1-20	1597		1994	0604			
	CA	2125359		AA	19941209		CA	1994	1-212	2535	9	1994	0607			
	AU	9464556		A1	19941215		AU	1994	1-64	556		1994	0607			
	AU	693786		В2	19980709											
	JP	07069894		A2	19950314		JP	1994	1-125	5091		1994	0607			
	FI	9402693		Α	19941209		FI	1994	1-269	93		1994	0608			
	NO	9402133		Α	19941209		NO	1994	1-213	33		1994	0608			
	HŲ	67413		A2	19950428		HU	1994	1-17	14		1994	0608			
	US	6121266		Α	20000919		US	1995	5-44	4984		1995	0519			
	US	6624166		В1	20030923		US	1996	5-62	5680		1996	0403			

- L9 ANSWER 8 OF 23 REGISTRY COPYRIGHT 2004 ACS on STN
- RN 161228-96-2 REGISTRY
- CN 4-Pyridazinecarboxamide, N-(2-amino-1-oxo-3-phenylpropyl)-3-chloro-6-(4-chlorophenyl)-, (S)- (9CI) (CA INDEX NAME)
- FS STEREOSEARCH
- MF C20 H16 C12 N4 O2

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

Absolute stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1: 122:160663 Preparation of 6-(hetero)aryl-3-chloroyridazine-4-carboxamides and analogs as interleukin-1β protease inhibitors.

Dolle, Roland E.; Hoyer, Denton W.; Schmidt, Stanley J.; Ross, Tina M.; Rinker, James M.; Ator, Mark A. (Sterling Winthrop Inc., USA). Eur. Pat. Appl. EP 628550 A2 19941214, 18 pp. DESIGNATED STATES: R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE. (English). CODEN: EPXXDW. APPLICATION: EP 1994-201597 19940604. PRIORITY: US 1993-73914 19930608.

	PA'	TENT NO.		KIND	DATE		APPLICATION NO.	DATE
PI	EP	628550		A2	19941214		EP 1994-201597	19940604
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	EP	628550		B1	19980225			
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	ES	2114654		Т3	19980601		ES 1994-201597	19940604
	CA	2125359		AA	19941209		CA 1994-2125359	19940607
	ΑU	9464556		A1	19941215		AU 1994-64556	19940607
	ΑU	693786		B2	19980709			
	JP	07069894		A2	19950314		JP 1994-125091	19940607
	FI	9402693		Α	19941209		FI 1994-2693	19940608
	NO	9402133		A	19941209		NO 1994-2133	19940608
	HU	67413		A2	19950428		HU 1994-1714	19940608
	US	6121266		Α	20000919		US 1995-444984	19950519
	US	6624166		B1	20030923		US 1996-625680	19960403

L9 ANSWER 9 OF 23 REGISTRY COPYRIGHT 2004 ACS on STN

RN 161228-95-1 REGISTRY

CN 4-Pyridazinecarboxamide, 3-chloro-6-[3-fluoro-5-methoxy-4-(methylsulfonyl)phenyl]- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C13 H11 C1 F N3 O4 S

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1: 122:160663 Preparation of 6-(hetero)aryl-3-chloroyridazine-4-carboxamides and analogs as interleukin-1β protease inhibitors.

Dolle, Roland E.; Hoyer, Denton W.; Schmidt, Stanley J.; Ross, Tina M.; Rinker, James M.; Ator, Mark A. (Sterling Winthrop Inc., USA). Eur. Pat. Appl. EP 628550 A2 19941214, 18 pp. DESIGNATED STATES: R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE. (English). CODEN: EPXXDW. APPLICATION: EP 1994-201597 19940604. PRIORITY: US 1993-73914 19930608.

	PAT	TENT NO.		KIND	DATE		APPLICATION NO.	DATE
PI		628550 628550		A2	19941214 19950118		EP 1994-201597	19940604
		628550		A3 B1	19980225			
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	ES	2114654		Т3	19980601		ES 1994-201597	19940604
	CA	2125359		AA	19941209		CA 1994-2125359	19940607
	ΑU	9464556		A 1	19941215		AU 1994-64556	19940607
	AU	693786		B2	19980709			
	JP	07069894		A2	19950314		JP 1994-125091	19940607
	FI	9402693		Α	19941209		FI 1994-2693	19940608
	ИО	9402133		Α	19941209		NO 1994-2133	19940608
	HU	67413		A 2	19950428		HU 1994-1714	19940608
	US	6121266		Α	20000919		US 1995-444984	19950519
	US	6624166		B1	20030923		US 1996-625680	19960403

L9 ANSWER 10 OF 23 REGISTRY COPYRIGHT 2004 ACS on STN

RN 161228-94-0 REGISTRY

CN 4-Pyridazinecarboxamide, 3-chloro-6-[3,5-difluoro-4-(methylsulfonyl)phenyl]- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C12 H8 C1 F2 N3 O3 S

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

$$\begin{array}{c|c} O & F \\ \parallel & & \\ \hline O & & \\ \hline & & \\ O & & \\ \hline & & \\ O & & \\ \hline & & \\ & & \\ & & \\ \hline & & \\ & & \\ & & \\ & & \\ \hline & & \\ & & \\ & & \\ & & \\ \hline & & \\ & & \\ & & \\ & & \\ \hline & & \\ & & \\ & & \\ & & \\ & & \\ \hline & & \\ & & \\ & & \\ & & \\ \hline & & \\ & & \\ & & \\ & & \\ & & \\ \end{array}$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 2 REFERENCES IN FILE CA (1907 TO DATE)
- 2 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- REFERENCE 1: 127:65731 3-Chloro-4-carboxamido-6-arylpyridazines as a
 non-peptide class of interleukin-1β converting enzyme inhibitor.
 Dolle, Roland E.; Hoyer, Denton; Rinker, James M.; Ross, Tina Morgan;
 Schmidt, Stanley J.; Helaszek, Carla; Ator, Mark A. (Sanofi Winthrop Inc.,
 Malvern, PA, 19426, USA). Bioorganic & Medicinal Chemistry Letters, 7(8),
 1003-1006 (English) 1997. CODEN: BMCLE8. ISSN: 0960-894X. Publisher:
 Elsevier.
- REFERENCE 2: 122:160663 Preparation of 6-(hetero)aryl-3-chloroyridazine-4-carboxamides and analogs as interleukin-1β protease inhibitors.

 Dolle, Roland E.; Hoyer, Denton W.; Schmidt, Stanley J.; Ross, Tina M.; Rinker, James M.; Ator, Mark A. (Sterling Winthrop Inc., USA). Eur. Pat. Appl. EP 628550 A2 19941214, 18 pp. DESIGNATED STATES: R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE. (English). CODEN: EPXXDW. APPLICATION: EP 1994-201597 19940604. PRIORITY: US 1993-73914 19930608.

	PAT	TENT NO.	KIND	DATE		APPLICATION NO.	DATE
ΡI	EP	628550	A2	19941214		EP 1994-201597	19940604
		628550	A3	19950118			
	EΡ	628550	B1	19980225			•
		R: AT, BE,	CH, DE	, DK, ES,	FR,	GB, GR, IE, IT, LI,	LU, MC, NL, PT, SE
	AT	163412	E	19980315		AT 1994-201597	19940604
	ES	2114654	Т3	19980601		ES 1994-201597	19940604
	CA	2125359	AA	19941209		CA 1994-2125359	19940607
	AU	9464556	A1	19941215		AU 1994-64556	19940607
	AU	693786	B2	19980709			
	JP	07069894	A2	19950314		JP 1994-125091	19940607
	FI	9402693	Α	19941209		FI 1994-2693	19940608
	NO	9402133	Α	19941209		NO 1994-2133	19940608
	HU	67413	A2	19950428		HU 1994-1714	19940608
	US	6121266	Α	20000919		US 1995-444984	19950519
	US	6624166	B1	20030923		US 1996-625680	19960403

- L9 ANSWER 11 OF 23 REGISTRY COPYRIGHT 2004 ACS on STN
- RN 161228-93-9 REGISTRY
- CN 4-Pyridazinecarboxamide, 3-chloro-6-(4-methoxyphenyl)- (9CI) (CA INDEX

NAME)

FS 3D CONCORD

MF C12 H10 C1 N3 O2

SR CF

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

$$C1$$
 $C-NH_2$
 O

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 2 REFERENCES IN FILE CA (1907 TO DATE)
- 2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1: 127:65731 3-Chloro-4-carboxamido-6-arylpyridazines as a
 non-peptide class of interleukin-1β converting enzyme inhibitor.
 Dolle, Roland E.; Hoyer, Denton; Rinker, James M.; Ross, Tina Morgan;
 Schmidt, Stanley J.; Helaszek, Carla; Ator, Mark A. (Sanofi Winthrop Inc.,
 Malvern, PA, 19426, USA). Bioorganic & Medicinal Chemistry Letters, 7(8),
 1003-1006 (English) 1997. CODEN: BMCLE8. ISSN: 0960-894X. Publisher:
 Elsevier.

	PAT	TENT NO.		KIND	DATE		API	PLICATI	ON NO.	DATE				
ΡI	EP	628550		A2	19941214		EP	1994-2	201597	1994	0604			
	ΕP	628550		A 3	19950118									
	EΡ	628550		B1	19980225									
		R: AT,	BE,	CH, DE	, DK, ES,	FR,	GB, G	GR, IE,	IT, L	I, LU,	MC,	NL,	PT,	SE
	AT	163412		E	19980315		AT	1994-2	201597	1994	0604			
	ES	2114654		Т3	19980601		ES	1994-2	201597	1994	0604			
	CA	2125359		AA	19941209		CA	1994 - 2	2125359	1994	0607			
	ΑU	9464556		A 1	19941215		AU	1994-6	54556	1994	0607			
	ΑU	693786		B2	19980709									
	JP	07069894		A2	19950314		JP	1994-1	L25091	1994	0607			
	FI	9402693		Α	19941209		FI	1994-2	2693	1994	0608			
	NO	9402133		Α	19941209		ИО	1994-2	2133	1994	0608			
	HU	67413		A2	19950428		HU	1994-1	1714	1994	0608			
	US	6121266		A	20000919		US	1995-4	144984	1995	0519			
	US	6624166		В1	20030923		US	1996-6	525680	1996	0403			

10/668,475 Thomas McKenzie

L9 ANSWER 12 OF 23 REGISTRY COPYRIGHT 2004 ACS on STN

RN 161228-91-7 REGISTRY

CN 4-Pyridazinecarboxamide, 3-chloro-6-[4-[[(4-methylphenyl)sulfonyl]amino]phenyl]- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C18 H15 C1 N4 O3 S

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1: 122:160663 Preparation of 6-(hetero)aryl-3-chloroyridazine-4-carboxamides and analogs as interleukin-1β protease inhibitors.

Dolle, Roland E.; Hoyer, Denton W.; Schmidt, Stanley J.; Ross, Tina M.; Rinker, James M.; Ator, Mark A. (Sterling Winthrop Inc., USA). Eur. Pat. Appl. EP 628550 A2 19941214, 18 pp. DESIGNATED STATES: R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE. (English). CODEN: EPXXDW. APPLICATION: EP 1994-201597 19940604. PRIORITY: US 1993-73914 19930608.

	PATENT NO.			KIND	DATE		API	PLICA	OITA	и ио.	DATE				
PI		628550 628550		A2 A3	19941214 19950118		EP	1994	4-201	1597	19940	0604			
		628550		В1	19980225										
		R: AT,	BE,	CH, DE	, DK, ES,	FR,	GB, G	GR, 1	IE, I	IT, LI	, LU,	MC,	NL,	PT,	SE
	ΑT	163412		E	19980315		ΑT	1994	4-20	1597	19940	0604			
	ES	2114654		Т3	19980601		ES	1994	4-20	1597	19940	0604			
	CA	2125359		AA	19941209		CA	1994	4-212	25359	19940	0607			
	ΑU	9464556		A1	19941215		AU	1994	4-645	556	19940	0607			
	ΑU	693786		B2	19980709										
	JP	07069894		A2	19950314		JP	1994	4-125	5091	19940	0607			
	FI	9402693		Α	19941209		FI	1994	4-269	93	19940	0608			
	NO	9402133		Α	19941209		ИО	1994	4-213	33	19940	8090			
	HU	67413		A2	19950428		HU	1994	4-17	14	19940	8090			
	US	6121266		A	20000919		US	1995	5-444	4984	19950)519			
	US	6624166		B1	20030923		US	1996	6-62	5680	19960	0403			

L9 ANSWER 13 OF 23 REGISTRY COPYRIGHT 2004 ACS on STN

RN 161228-87-1 REGISTRY

CN 4-Pyridazinecarboxamide, 3-chloro-6-(4-chlorophenyl)-5-methyl- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C12 H9 C12 N3 O

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

$$\begin{array}{c|c} N & N & \\ \hline N & \\ C & \\ C - NH_2 & \\ 0 & \\ \end{array}$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 2 REFERENCES IN FILE CA (1907 TO DATE)
 2 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- REFERENCE 1: 127:65731 3-Chloro-4-carboxamido-6-arylpyridazines as a
 non-peptide class of interleukin-1β converting enzyme inhibitor.
 Dolle, Roland E.; Hoyer, Denton; Rinker, James M.; Ross, Tina Morgan;
 Schmidt, Stanley J.; Helaszek, Carla; Ator, Mark A. (Sanofi Winthrop Inc.,
 Malvern, PA, 19426, USA). Bioorganic & Medicinal Chemistry Letters, 7(8),
 1003-1006 (English) 1997. CODEN: BMCLE8. ISSN: 0960-894X. Publisher:
 Elsevier.
- REFERENCE 2: 122:160663 Preparation of 6-(hetero)aryl-3-chloroyridazine-4-carboxamides and analogs as interleukin-1β protease inhibitors.

 Dolle, Roland E.; Hoyer, Denton W.; Schmidt, Stanley J.; Ross, Tina M.; Rinker, James M.; Ator, Mark A. (Sterling Winthrop Inc., USA). Eur. Pat. Appl. EP 628550 A2 19941214, 18 pp. DESIGNATED STATES: R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE. (English).

 CODEN: EPXXDW. APPLICATION: EP 1994-201597 19940604. PRIORITY: US 1993-73914 19930608.

	PATENT NO.			KIND	DATE		APPLICATION NO.	DATE
ΡI	EΡ	628550		A2	19941214		EP 1994-201597	19940604
	EΡ	628550		A3	19950118			
	EΡ	628550		В1	19980225			
		R: AT,	BE,	CH, DE	, DK, ES,	FR,	GB, GR, IE, IT, LI,	LU, MC, NL, PT, SE
	AT	163412		E	19980315		AT 1994-201597	19940604
	ES	2114654		Т3	19980601		ES 1994-201597	19940604
	CA	2125359		AA	19941209		CA 1994-2125359	19940607
	AU	9464556		A1	19941215		AU 1994-64556	19940607
	AU	693786		В2	19980709			
	JP	07069894		A2	19950314		JP 1994-125091	19940607
	FI	9402693		A	19941209		FI 1994-2693	19940608
	NO	9402133		Α	19941209		NO 1994-2133	19940608
	HU	67413		A2	19950428		HU 1994-1714	19940608
	US	6121266		Α	20000919		US 1995-444984	19950519
	US	6624166		В1	20030923		US 1996-625680	19960403

L9 ANSWER 14 OF 23 REGISTRY COPYRIGHT 2004 ACS on STN RN 161228-85-9 REGISTRY

4-Pyridazinecarboxamide, 3-chloro-6-(3-chloro-4-cyanophenyl)- (9CI) CN (CA INDEX NAME)

FS 3D CONCORD

C12 H6 C12 N4 O MF

SR

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 2 REFERENCES IN FILE CA (1907 TO DATE)
- 2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1: 127:65731 3-Chloro-4-carboxamido-6-arylpyridazines as a non-peptide class of interleukin- 1β converting enzyme inhibitor. Dolle, Roland E.; Hoyer, Denton; Rinker, James M.; Ross, Tina Morgan; Schmidt, Stanley J.; Helaszek, Carla; Ator, Mark A. (Sanofi Winthrop Inc., Malvern, PA, 19426, USA). Bioorganic & Medicinal Chemistry Letters, 7(8), 1003-1006 (English) 1997. CODEN: BMCLE8. ISSN: 0960-894X. Publisher: Elsevier.

	PA'	CENT NO.		KIND	DATE		API	PLICATI	ON NO).	DATE				
PI	EP	628550		A2	19941214		EP	1994-2	201597	7	19940	604			
	EP	628550		A3	19950118										
	EP	628550		В1	19980225										
		R: AT,	BE,	CH, DE	, DK, ES,	FR,	GB, G	SR, IE,	IT,	LI,	LU,	MC,	NL,	PT,	SE
	ΑT	163412		E	19980315		AΤ	1994-2	201597	7	19940	604			
	ES	2114654		Т3	19980601		ES	1994-2	201597	7	19940	604			
	CA	2125359		AA	19941209		CA	1994-2	212535	59	19940	607			
	AU	9464556		A1	19941215		AU	1994-6	54556		19940	607			
	AU	693786		B2	19980709										
	JP	07069894		A2	19950314		JP	1994-1	25091	L	19940	607			
	FΙ	9402693		Α	19941209		FI	1994-2	2693		19940	608			
	NO	9402133		Α	19941209		NO	1994-2	2133		19940	608			
	ΗU	67413		A2	19950428		HU	1994-1	714		19940	608			
	US	6121266		Α	20000919		US	1995-4	144984	1	19950	519			
	US	6624166		В1	20030923		US	1996-6	25680)	19960	403			

L9 ANSWER 15 OF 23 REGISTRY COPYRIGHT 2004 ACS on STN

RN 161228-84-8 REGISTRY

CN 4-Pyridazinecarboxamide, 3-chloro-6-(4-nitrophenyl)- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C11 H7 C1 N4 O3

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1907 TO DATE)
2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1: 127:65731 3-Chloro-4-carboxamido-6-arylpyridazines as a non-peptide class of interleukin-1β converting enzyme inhibitor.

Dolle, Roland E.; Hoyer, Denton; Rinker, James M.; Ross, Tina Morgan; Schmidt, Stanley J.; Helaszek, Carla; Ator, Mark A. (Sanofi Winthrop Inc., Malvern, PA, 19426, USA). Bioorganic & Medicinal Chemistry Letters, 7(8), 1003-1006 (English) 1997. CODEN: BMCLE8. ISSN: 0960-894X. Publisher: Elsevier.

PATENT NO.				KIN	ID	DATE			AF	PLI	CATIO	ои ис).	DATE				
EΡ	62855	50		A3	}	1995	0118		EF	19	94-20	01597	7	1994	0604			
ĿĘ			BE,		_			FR,	GB,	GR,	IE,	IT,	LI,	LU,	MC,	NL,	PT,	SE
ΑT	16341	12		Ë		1998	0315		ĀT	19	94-20	01597	7	1994	0604			
ES	21146	654		Т3	}	1998	0601		ES	19	94-20	01597	7	1994	0604			
CA	21253	359		A.F	A.	1994	1209		C.P	. 19	94-2	12535	59	1994	0607			
AU	94645	556		A1	_	1994	1215		AU	19	94-64	4556		1994	0607			
UA	69378	36		B2	?	1998	0709											
JP	07069	9894		A2	·	1995	0314		JE	19	94-12	25093	L	1994	0607			
FΙ	94026	593		Α		1994	1209		FI	19	94-2	693		1994	0608			
ИО	94021	133		Α		1994	1209		NC	19	94-2	133		1994	0608			
	EP EP EP AT ES AU AU UP	EP 62855 EP 62855 R: AT 16341 ES 21146 CA 21253 AU 94645 AU 69376 JP 07066 FI 94026	EP 628550 EP 628550 EP 628550	EP 628550 EP 628550 EP 628550 R: AT, BE, AT 163412 ES 2114654 CA 2125359 AU 9464556 AU 693786 JP 07069894 FI 9402693	EP 628550 A2 EP 628550 B1 EP 628550 B1 R: AT, BE, CH, AT 163412 E ES 2114654 T3 CA 2125359 AA AU 9464556 A1 AU 693786 B2 JP 07069894 A2 FI 9402693 A	EP 628550 A2 EP 628550 B1 R: AT, BE, CH, DE, AT 163412 E ES 2114654 T3 CA 2125359 AA AU 9464556 A1 AU 693786 B2 JP 07069894 A2 FI 9402693 A	EP 628550 A2 1994 EP 628550 B1 1998 EP 628550 B1 1998 EP 628550 B1 1998 AT 163412 E 1998 ES 2114654 T3 1998 AD 2125359 AA 1994 AU 9464556 A1 1994 AU 693786 B2 1998 AU 693786 B2 1998 AU 69402693 A 1994	EP 628550 A2 19941214 EP 628550 B1 19980225 R: AT, BE, CH, DE, DK, ES, AT 163412 E 19980315 ES 2114654 T3 19980601 CA 2125359 AA 19941209 AU 9464556 A1 19941215 AU 693786 B2 19980709 JP 07069894 A2 19950314 EI 9402693 A 19941209	EP 628550 A2 19941214 EP 628550 B1 19980225 EP 628550 B1 19980315 EP 628550 B2 19980601 EP 693786 B2 19980709 EP 693786 B2 19950314 EP 628550 A3 19941209	EP 628550 A2 19941214 EP EP 628550 B1 19980225 R: AT, BE, CH, DE, DK, ES, FR, GB, AT 163412 E 19980315 AT ES 2114654 T3 19980601 ES CA 2125359 AA 19941209 CA AU 9464556 A1 19941215 AU AU 693786 B2 19980709 JP 07069894 A2 19950314 JP EF 9402693 A 19941209 FI	EP 628550 A2 19941214 EP 19 EP 628550 B1 19980225 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, AT 163412 E 19980315 AT 19 ES 2114654 T3 19980601 ES 19 CA 2125359 AA 19941209 CA 19 AU 9464556 A1 19941215 AU 19 AU 693786 B2 19980709 JP 07069894 A2 19950314 JP 19 EF 9402693 A 19941209 FI 19	EP 628550 A2 19941214 EP 1994-20 EP 628550 B1 19980225 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, AT 163412 E 19980315 AT 1994-20 ES 2114654 T3 19980601 ES 1994-20 CA 2125359 AA 19941209 CA 1994-20 AU 9464556 A1 19941215 AU 1994-60 AU 693786 B2 19980709 JP 07069894 A2 19950314 JP 1994-12 ET 9402693 A 19941209 FI 1994-20	EP 628550 A2 19941214 EP 1994-20159 EP 628550 B1 19980225 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, AT 163412 E 19980315 AT 1994-20159 ES 2114654 T3 19980601 ES 1994-20159 CA 2125359 AA 19941209 CA 1994-212535 AU 9464556 A1 19941215 AU 1994-64556 AU 693786 B2 19980709 JP 07069894 A2 19950314 JP 1994-125093 ET 9402693 A 19941209 FI 1994-2693	EP 628550 A2 19941214 EP 1994-201597 EP 628550 B1 19980225 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, AT 163412 E 19980315 AT 1994-201597 ES 2114654 T3 19980601 ES 1994-201597 CA 2125359 AA 19941209 CA 1994-2125359 AU 9464556 A1 19941215 AU 1994-64556 AU 693786 B2 19980709 JP 07069894 A2 19950314 JP 1994-125091 FI 9402693 A 19941209 FI 1994-2693	EP 628550 A2 19941214 EP 1994-201597 1994 EP 628550 B1 19980225 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, AT 163412 E 19980315 AT 1994-201597 1994 ES 2114654 T3 19980601 ES 1994-201597 1994 CA 2125359 AA 19941209 CA 1994-2125359 1994 AU 9464556 A1 19941215 AU 1994-64556 1994 AU 693786 B2 19980709 JP 07069894 A2 19950314 JP 1994-125091 1994 FI 9402693 A 19941209 FI 1994-2693 1994	EP 628550 A2 19941214 EP 1994-201597 19940604 EP 628550 B1 19980225 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, AT 163412 E 19980315 AT 1994-201597 19940604 ES 2114654 T3 19980601 ES 1994-201597 19940604 EA 2125359 AA 19941209 CA 1994-2125359 19940607 EAU 9464556 A1 19941215 AU 1994-64556 19940607 EAU 693786 B2 19980709 ED 07069894 A2 19950314 JP 1994-125091 19940607 ET 9402693 A 19941209 FI 1994-2693 19940608	EP 628550 A2 19941214 EP 1994-201597 19940604 EP 628550 B1 19980225 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, AT 163412 E 19980315 AT 1994-201597 19940604 ES 2114654 T3 19980601 ES 1994-201597 19940604 EA 2125359 AA 19941209 CA 1994-2125359 19940607 EAU 946456 A1 19941215 AU 1994-64556 19940607 EAU 693786 B2 19980709 ED 07069894 A2 19950314 JP 1994-125091 19940607 ET 9402693 A 19941209 FI 1994-2693 19940608	EP 628550 A2 19941214 EP 1994-201597 19940604 EP 628550 B1 19980225 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, AT 163412 E 19980315 AT 1994-201597 19940604 ES 2114654 T3 19980601 ES 1994-201597 19940604 EA 2125359 AA 19941209 CA 1994-2125359 19940607 EAU 946456 A1 19941215 AU 1994-64556 19940607 EU 9402693 A 19941209 FI 1994-2693 19940608

10/668,475 Thomas McKenzie

HU 67413	A2	19950428	HU 1994-1714	19940608
US 6121266	Α	20000919	US 1995-444984	19950519
US 6624166	В1	20030923	US 1996-625680	19960403

L9 ANSWER 16 OF 23 REGISTRY COPYRIGHT 2004 ACS on STN

RN 161228-82-6 REGISTRY

CN 4-Pyridazinecarboxamide, 3-chloro-6-(3,5-dichlorophenyl)- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C11 H6 C13 N3 O

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

$$\begin{array}{c|c} C1 & N & N \\ \hline & C1 & C-NH_2 \\ \hline & O & \end{array}$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1: 122:160663 Preparation of 6-(hetero)aryl-3-chloroyridazine-4-carboxamides and analogs as interleukin-1β protease inhibitors.

Dolle, Roland E.; Hoyer, Denton W.; Schmidt, Stanley J.; Ross, Tina M.; Rinker, James M.; Ator, Mark A. (Sterling Winthrop Inc., USA). Eur. Pat. Appl. EP 628550 A2 19941214, 18 pp. DESIGNATED STATES: R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE. (English).

CODEN: EPXXDW. APPLICATION: EP 1994-201597 19940604. PRIORITY: US 1993-73914 19930608.

	PATENT NO.			KIND	DATE		API	PLICATI	ON NO	. DA	TE			
ΡI	EP	628550		A2	19941214		EP	1994-2	 201597	19	940604			
	EP	628550		A3	19950118									
	EP	628550		B1	19980225									
		R: AT,	BE,	CH, DE,	, DK, ES,	FR,	GB, G	R, IE,	IT, 1	LI, I	U, MC,	NL,	PT,	SE
	AT	163412		E	19980315		AT	1994-2	201597	19	940604			
	ES	2114654		Т3	19980601		ES	1994-2	201597	19	940604			
	CA	2125359		AA	19941209		CA	1994-2	2125359	9 19	940607			
	AU	9464556		A1	19941215		AU	1994-6	54556	19	940607			
	AU	693786		B2	19980709									
	JP	07069894		A2	19950314		JP	1994-1	L25091	19	940607			
	FI	9402693		Α	19941209		FI	1994-2	2693	19	940608			
	NO	9402133		Α	19941209		NO	1994-2	2133	19	940608			
	HU	67413		A2	19950428		HU	1994-1	L714	19	940608			
	US	6121266		Α	20000919		US	1995-4	144984	19	950519			
	US	6624166		B1	20030923		US	1996-6	525680	19	960403			

L9 ANSWER 17 OF 23 REGISTRY COPYRIGHT 2004 ACS on STN

RN 161228-81-5 REGISTRY

CN 4-Pyridazinecarboxamide, 3-chloro-6-[4-(trifluoromethyl)phenyl]- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C12 H7 C1 F3 N3 O

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

$$C1$$
 CF_3
 $C = NH_2$
 $C = NH_2$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1907 TO DATE)
2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1: 127:65731 3-Chloro-4-carboxamido-6-arylpyridazines as a non-peptide class of interleukin-1β converting enzyme inhibitor.

Dolle, Roland E.; Hoyer, Denton; Rinker, James M.; Ross, Tina Morgan; Schmidt, Stanley J.; Helaszek, Carla; Ator, Mark A. (Sanofi Winthrop Inc., Malvern, PA, 19426, USA). Bioorganic & Medicinal Chemistry Letters, 7(8), 1003-1006 (English) 1997. CODEN: BMCLE8. ISSN: 0960-894X. Publisher: Elsevier.

	PAT	TENT NO.		KIND	DATE		APPLICATION NO		DATE			
								-				
ΡI	EP	628550		A2	19941214		EP 1994-201597	,	19940604			
	EP	628550		A 3	19950118							
	EP	628550		B1	19980225							
		R: AT,	BE,	CH, DE	, DK, ES,	FR,	GB, GR, IE, IT,	LI,	LU, MC,	NL,	PT,	SE
	AT	163412		E	19980315		AT 1994-201597		19940604			
	ES	2114654		Т3	19980601		ES 1994-201597	•	19940604			
	CA	2125359		AA	19941209		CA 1994-212535	9	19940607			
	ΑU	9464556		A 1	19941215		AU 1994-64556		19940607			
	ΑU	693786		B2	19980709							
	JP	07069894		A2	19950314		JP 1994-125091	_	19940607			
	FI	9402693		Α	19941209		FI 1994-2693		19940608			
	NO	9402133		Α	19941209		NO 1994-2133		19940608			
	HU	67413		A2	19950428		HU 1994-1714		19940608			
	US	6121266		Α	20000919		US 1995-444984		19950519			

US 6624166 B1 20030923 US 1996-625680 19960403

L9 ANSWER 18 OF 23 REGISTRY COPYRIGHT 2004 ACS on STN

RN 161228-80-4 REGISTRY

CN 4-Pyridazinecarboxamide, 6-(4-bromophenyl)-3-chloro- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C11 H7 Br Cl N3 O

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1907 TO DATE)
2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1: 127:65731 3-Chloro-4-carboxamido-6-arylpyridazines as a non-peptide class of interleukin-1β converting enzyme inhibitor. Dolle, Roland E.; Hoyer, Denton; Rinker, James M.; Ross, Tina Morgan; Schmidt, Stanley J.; Helaszek, Carla; Ator, Mark A. (Sanofi Winthrop Inc., Malvern, PA, 19426, USA). Bioorganic & Medicinal Chemistry Letters, 7(8), 1003-1006 (English) 1997. CODEN: BMCLE8. ISSN: 0960-894X. Publisher: Elsevier.

	PAT	CENT	NO.		KIN	ID	DATE			ΑE	bri	CATIO	ON NO).	DATE				
PI	ΕP	6285	550		A2	2	1994	1214		EF	19	94-20	1597	7	1994	0604			
	ΕP	6285	550		A3	3	1995	0118											
	EΡ	P 628550 R: AT, BE,			В1	_	1998	0225											
		R:	AT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	ΙE,	IT,	LI,	LU,	MC,	NL,	PT,	SE
	ΑT	1634	112		E		1998	0315		ΓA	19	94-20	01597	7	1994	0604			
	ES	2114	1654		Т3	}	1998	0601		ES	19	94-20	01597	7	1994	0604			
	CA	2125	5359		AA	A.	1994	1209		CF	19	94-23	12535	59	1994	0607			
	AU	9464	1556		A1		1994	1215		ΑU	J 19	94-64	1556		1994	0607			
	AU	6937	786		B2	2	1998	0709		-									
	JP	0706	59894		A2	?	1995	0314		JE	19	94-12	25091	L	1994	0607			
	FI	9402	2693		Α		1994	1209		FI	19	94-26	593		1994	0608			

NO 9402133	Α	19941209	NO :	1994-2133	19940608
HU 67413 /	A2	19950428	HU	1994-1714	19940608
US 6121266	Α	20000919	US :	1995-444984	19950519
(US 6624166)	B1	20030923	US :	1996-625680	19960403

L9 ANSWER 19 OF 23 REGISTRY COPYRIGHT 2004 ACS on STN

RN 150594-44-8 REGISTRY

CN Cyclohexanecarboxylic acid, 4-[[[3-chloro-6-(4-cyanophenyl)-4-pyridazinyl]carbonyl]amino]-, methyl ester, trans- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C20 H19 C1 N4 O3

DATENT NO

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

Relative stereochemistry.

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1: 119:225966 Preparative and biological activity of aryl substituted nitrogen containing heterocycles. Linz, Guenter; Pieper, Helmut; Himmelsbach, Frank; Austel, Volkhard; Mueller, Thomas; Weisenberger, Johannes; Seewaldt-Becker, Elke (Thomae, Dr. Karl, G.m.b.H., Germany). Eur. Pat. Appl. EP 537696 Al 19930421, 47 pp. DESIGNATED STATES: R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, NL, PT, SE. (German). CODEN: EPXXDW. APPLICATION: EP 1992-117507 19921014. PRIORITY: DE 1991-4134467 19911018.

ADDITCATION NO

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	PAIENI NO.	KIND DAIE	APPLICATION NO.	DAIL
ΡI	EP 537696	A1 1993042	EP 1992-117507	19921014
	R: AT, BE,	CH, DE, DK, ES	FR, GB, GR, IE, IT, LI,	LU, NL, PT, SE
	DE 4134467	A1 1993042	DE 1991-4134467	19911018
	US 5418233	A 1995052	3 US 1992-961135	19921014
	CA 2080748	AA 1993041	CA 1992-2080748	19921016
	NO 9204027	A 1993041	NO 1992-4027	19921016
	AU 9227062	A1 1993042	2 AU 1992-27062	19921016
	AU 662930	B2 1995092	L	
	HU 62272	A2 1993042	B HU 1992-3264	19921016
	JP 05221992	A2 1993083	l JP 1992-277578	19921016
	ZA 9207994	A 1994041	3 ZA 1992-7994	19921016
	US 5563268	A 1996100	3 US 1995-375084	19950119

L9 ANSWER 20 OF 23 REGISTRY COPYRIGHT 2004 ACS on STN

KIND DATE

10/668,475 Thomas McKenzie

RN 150594-34-6 REGISTRY

CN 4-Pyridazinecarboxylic acid, 3-chloro-6-(4-cyanophenyl)- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C12 H6 C1 N3 O2

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1: 119:225966 Preparative and biological activity of aryl substituted nitrogen containing heterocycles. Linz, Guenter; Pieper, Helmut; Himmelsbach, Frank; Austel, Volkhard; Mueller, Thomas; Weisenberger, Johannes; Seewaldt-Becker, Elke (Thomae, Dr. Karl, G.m.b.H., Germany). Eur. Pat. Appl. EP 537696 A1 19930421, 47 pp. DESIGNATED STATES: R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, NL, PT, SE. (German). CODEN: EPXXDW. APPLICATION: EP 1992-117507 19921014. PRIORITY: DE 1991-4134467 19911018.

	PATENT NO.	KIND DATE	APPLICATION NO. DATE	
ΡI	EP 537696	A1 19930421	EP 1992-117507 19921014	
	R: AT, BE,	CH, DE, DK, ES, FR,	GB, GR, IE, IT, LI, LU, NL, PT	, SE
	DE 4134467	Al 19930422	DE 1991-4134467 19911018	
	US 5418233	A 19950523	US 1992-961135 19921014	
	CA 2080748	AA 19930419	CA 1992-2080748 19921016	
	NO 9204027	A 19930419	NO 1992-4027 19921016	
	AU 9227062	A1 19930422	AU 1992-27062 19921016	
	AU 662930	B2 19950921		
	HU 62272	A2 19930428	HU 1992-3264 19921016	
	JP 05221992	A2 19930831	JP 1992-277578 19921016	
	ZA 9207994	A 19940418	ZA 1992-7994 19921016	
	US 5563268	A 19961008	US 1995-375084 19950119	

L9 ANSWER 21 OF 23 REGISTRY COPYRIGHT 2004 ACS on STN

RN 150594-29-9 REGISTRY

CN 4-Pyridazinecarboxylic acid, 3-chloro-6-(4-cyanophenyl)-, ethyl ester (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C14 H10 Cl N3 O2

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 1 REFERENCES IN FILE CA (1907 TO DATE)
- 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1: 119:225966 Preparative and biological activity of aryl substituted nitrogen containing heterocycles. Linz, Guenter; Pieper, Helmut; Himmelsbach, Frank; Austel, Volkhard; Mueller, Thomas; Weisenberger, Johannes; Seewaldt-Becker, Elke (Thomae, Dr. Karl, G.m.b.H., Germany). Eur. Pat. Appl. EP 537696 A1 19930421, 47 pp. DESIGNATED STATES: R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, NL, PT, SE. (German). CODEN: EPXXDW. APPLICATION: EP 1992-117507 19921014. PRIORITY: DE 1991-4134467 19911018.

PAT	TENT NO.	KIND	DATE	APPLICATION NO. DATE	
PI EP	537696	A1	19930421	EP 1992-117507 19921014	
	R: AT, BE,	CH, DE,	DK, ES,	FR, GB, GR, IE, IT, LI, LU, NL, PT, SE	
DE	4134467	A1	19930422	DE 1991-4134467 19911018	
US	5418233	A	19950523	US 1992-961135 19921014	
CA	2080748	AA	19930419	CA 1992-2080748 19921016	
NO	9204027	Α	19930419	NO 1992-4027 19921016	
AU	9227062	A1	19930422	AU 1992-27062 19921016	
AU	662930	B2	19950921		
HU	62272	A2	19930428	ни 1992-3264 19921016	
JP	05221992	A 2	19930831	JP 1992-277578 19921016	
ZA	9207994	Α	19940418	ZA 1992-7994 19921016	
US	5563268	Α	19961008	US 1995-375084 19950119	

- L9 ANSWER 22 OF 23 REGISTRY COPYRIGHT 2004 ACS on STN
- RN 101884-08-6 REGISTRY
- CN 4-Pyridazinecarboxylic acid, 3-chloro-5,6-bis(p-chlorophenyl)-, ethyl ester (6CI) (CA INDEX NAME)
- FS 3D CONCORD
- MF C19 H13 C13 N2 O2
- SR CAOLD
- LC STN Files: CA, CAOLD, CAPLUS

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

REFERENCE 1: 53:122325 6-Chloropyridazines. (C I B A Ltd.). GB 807548

19590114 (Unavailable). APPLICATION: GB.

PATENT NO. KIND DATE APPLICATION NO. DATE

PI GB 807548 19590114 GB

L9 ANSWER 23 OF 23 REGISTRY COPYRIGHT 2004 ACS on STN

RN 70769-18-5 REGISTRY

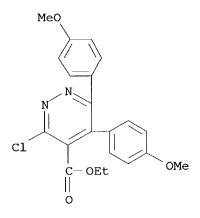
CN 4-Pyridazinecarboxylic acid, 3-chloro-5,6-bis(4-methoxyphenyl)-, ethyl ester (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C21 H19 C1 N2 O4

LC STN Files: BEILSTEIN*, CA, CAPLUS, CASREACT

(*File contains numerically searchable property data)



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 2 REFERENCES IN FILE CA (1907 TO DATE)
 2 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- REFERENCE 1: 118:213015 Heterocyclic β -enamino esters. 54. A new, general approach for the synthesis of heteroannulated 3,1-oxazin-4-ones. Wamhoff, Heinrich; Herrmann, Stefan; Stoelben, Stephan; Nieger, Martin (Inst. Org. Chem. Biochem., Univ. Bonn, Bonn, D-5300, Germany). Tetrahedron, 49(3), 581-94 (English) 1993. CODEN: TETRAB. ISSN: 0040-4020.
- REFERENCE 2: 91:56936 Synthesis and pharmacological activity of some 5,6-diphenyl-pyridazines. Nannini, Giuliano; Biasoli, Giovanni; Perrone, Ettore; Forgione, Angelo; Buttinoni, Ada; Ferrari, Mario (Carlo Erba Res. Inst., Milan, 20159, Italy). European Journal of Medicinal Chemistry, 14(1), 53-60 (English) 1979. CODEN: EJMCA5. ISSN: 0009-4374.

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ALL L# QUERIES AND ANSWER SETS ARE DELETED AT LOGOFF
LOGOFF? (Y)/N/HOLD:.
STN INTERNATIONAL LOGOFF AT 13:16:40 ON 27 FEB 2004